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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
09/665,472	09/20/2000	Menzo Havenga	4489US	8505
24247	7590 09/15/2003			
TRASK BRITT			EXAMINER	
P.O. BOX 2550 SALT LAKE CITY, UT 84110			MARVICH, MARIA	
			ART UNIT	PAPER NUMBER
			1636	
			DATE MAILED: 09/15/2003	

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)			
Advisory Action	09/665,472	HAVENGA ET AL.			
Advisory Action	Examiner	Art Unit			
	Maria B Marvich, PhD	1636			
Th MAILING DATE of this communication appe	ars on the cover sheet with the c	orrespondence address			
THE REPLY FILED 22 August 2003 FAILS TO PLACE Therefore, further action by the applicant is required to a final rejection under 37 CFR 1.113 may only be either: (1 condition for allowance; (2) a timely filed Notice of Appearance (RCE) in compliance with 37 CFR 1.114.	void abandonment of this application in the same of th	cation. A proper reply to a chapter chapter chapter the application in			
PERIOD FOR RE	PLY [check either a) or b)]				
a) The period for reply expires 3 months from the mailing date of b) The period for reply expires on: (1) the mailing date of this Adviewent, however, will the statutory period for reply expire later the ONLY CHECK THIS BOX WHEN THE FIRST REPLY WAS 706.07(f).	isory Action, or (2) the date set forth in the an SIX MONTHS from the mailing date of	f the final rejection.			
Extensions of time may be obtained under 37 CFR 1.136(a). The dat have been filed is the date for purposes of determining the period of extens 37 CFR 1.17(a) is calculated from: (1) the expiration date of the shortened (b) above, if checked. Any reply received by the Office later than three more earned patent term adjustment. See 37 CFR 1.704(b).	sion and the corresponding amount of the I statutory period for reply originally set in t	fee. The appropriate extension fee under the final Office action; or (2) as set forth in			
1. A Notice of Appeal was filed on Appellant's 37 CFR 1.192(a), or any extension thereof (37 CFR					
2. The proposed amendment(s) will not be entered be	ecause:				
(a) $oxed{\boxtimes}$ they raise new issues that would require furthe	er consideration and/or search (see NOTE below);			
(b) \square they raise the issue of new matter (see Note b	pelow);				
(c)	n better form for appeal by mate	erially reducing or simplifying the			
(d) They present additional claims without cancel	ing a corresponding number of f	finally rejected claims.			
NOTE: See Continuation Sheet.					
3. Applicant's reply has overcome the following reject	tion(s):				
4. Newly proposed or amended claim(s) would canceling the non-allowable claim(s).	be allowable if submitted in a se	eparate, timely filed amendment			
5. ☐ The a) ☐ affidavit, b) ☐ exhibit, or c) ☐ request for application in condition for allowance because: Se		idered but does NOT place the			
6. The affidavit or exhibit will NOT be considered bed raised by the Examiner in the final rejection.	cause it is not directed SOLELY	to issues which were newly			
7. For purposes of Appeal, the proposed amendment(s) a) will not be entered or b) will be entered and an explanation of how the new or amended claims would be rejected is provided below or appended.					
The status of the claim(s) is (or will be) as follows:					
Claim(s) allowed:					
Claim(s) objected to:		11			
Claim(s) rejected: <u>1,4,7,8,13,14,16,18,19 and 21-24</u> .					
Claim(s) withdrawn from consideration:					
8. The proposed drawing correction filed on is	a) approved or b) disapp	proved by the Examiner.			
9. Note the attached Information Disclosure Statemen	nt(s)(PTO-1449) Paper No(s)				
10. Other:		Linga Mileley			
•		TERRY MCKELVEY PRIMARY EXAMINER			

Continuation Sheet (PTOL-303) 009/665,472

Application No.

Continuation of 2. NOTE: The proposed amendment would raise the following new issues that would require new consideration: Claim 1 recites that the recombinant chimeric adenoviral vector is based on a first adenovirus of subgroup C and that the tissue tropism for dendritic cells is provided by a non-native fiber protein substituted for a fiber protein of the first adenovirus. These amendments have changed the scope of the claims by specifically limiting the vector as based on adenovirus of subgroup C whose native fiber protein is substituted with a non-native fiber from adenvoirus 11, 16, 35, 51 and 40L.

Continuation of 5. does NOT place the application in condition for allowance because: the applicant's arguments are most in view of the non-entry of the after final amendment.

On pages 7-8 of the amendment filed 12/17/02, applicant traverses the rejection of claims 1, 4, 7-8, 13-14, 16, 18, 19 and 21-22 under 35 U.S.C 102(e) as anticipated by Crystal et al. Applicant argues that Crytal et al neither expressly nor inherently teach the instant invention. Specifically, Crystal et al do not provide for modifications in adenoviral fiber proteins directed to provide tropism to target cells and specifically to dendritic cells. Instead, the reference to the teachings of Crystal et al is disputed as broadly describing all chimeric adenoviral vectors with switched fiber proteins. Teachings of modifications directed to altered tropism specifically for dendritic cells are said to be are attributed to Crystal et al based upon the theory of inherency. While according to the MPEP 2112, the examiner must provide a basis in fact or technical reasoning to demonstrate that the inherent characteristics necessarily flows from the teachings of the prior art, support for inherency is drawn from the specification of the instant invention. Applicants argue that the instant invention is distinguishable over the prior art by teaching that selected adenoviral serotypes or groups of serotypes with particularily advantageous tropism for dendritic cells. Specifically, they teach that not all adenovirus have tropism for dendritic cells and rather the art of selecting adenoviral fibers for tropism to dendritic cells as well as the tropism of each serotype is unpredictable. Further these teachings of the specification provide evidence that chimeric fibers made according to the teachings of Crystal et al would not possess the ability to bind and infect dendritic cells.

Applicants arguments filed 8/22/03 have been considered but are not persuasive. The elements disputed as missing from the teachings of Crystal et al, modifications directed to providing tropism to desired target cells especially dendritic cells, attribute to the claim processes that are not recited in the claim. Instead, applicants recite a chimeric adenoviral vector comprised of a first adenovirus capsid from Ad11, Ad 16, Ad35, Ad51, Ad40L and a second capsid from subgroup C adenovirus. The first capsid directs tropism for dendritic cells. This product is identical to that of Crystal et al. Therefore, the claimed properties are presumed to be inherent (see MPEP 2112.01). "Where the claimed and prior art products are identical or substantially identical in structure or composition, or are produced by identical or substantially identical processes, a prima facie case of either anticipation or obviousness has been established". Applicants have not provided evidence showing the prior art products do not necessarly possess the characteristics of the claimed product. As the prior art adenoviral vector and the adenoviral vector if the instant invention are identical and these facts are not in dispute, it would necessarily flow that the properties associated with these vectors are the same.